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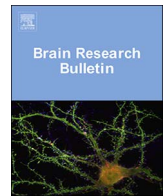
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Research report

Age-dependent changes in autophosphorylation of alpha calcium/calmodulin dependent kinase II in hippocampus and amygdala after contextual fear conditioning



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ABSTRACT

The hippocampus and amygdala are essential brain regions responsible for contextual fear conditioning (CFC). The autophosphorylation of alpha calcium-calmodulin kinase II (α CaMKII) at threonine-286 (T286) is a critical step implicated in long-term potentiation (LTP), learning and memory. However, the changes in α CaMKII levels with aging and training in associated brain regions are not fully understood. Here, we studied how aging and training affect the levels of phosphorylated (T286) and proportion of phosphorylated:total α CaMKII in the hippocampus and amygdala. Young and aged mice, naïve (untrained) and trained in CFC, were analysed by immunohistochemistry for the levels of total and phosphorylated α CaMKII in the hippocampus and amygdala. We found that two hours after CFC training, young mice exhibited a higher level of phosphorylated and increased ratio of phosphorylated:total α CaMKII in hippocampal CA3 stratum radiatum. Furthermore, aged untrained mice showed a higher ratio of phosphorylated:total α CaMKII in the CA3 region of the hippocampus when compared to the young untrained group. No effect of training or aging were seen in the central, lateral and basolateral amygdala regions, for both phosphorylated and ratio of phosphorylated:total α CaMKII. These results show that aging impairs the training-induced upregulation of autophosphorylated (T286) α CaMKII in the CA3 stratum radiatum of the hippocampus. This indicates that distinct age-related mechanisms underlie CFC that may rely more heavily on NMDA receptor-dependent plasticity in young age.

1. Introduction

In the hippocampus, long-term potentiation (LTP) has been implicated as one of the memory mechanisms (Giese, 2012). Autophosphorylation at the threonine-286 (T286) of alpha calcium/calmodulin-dependent kinase II (α CaMKII) is critical to inducing NMDA receptor-dependent LTP in hippocampal area CA1 (Lisman et al., 2012; Giese and Mizuno, 2013). The absence of this LTP combined with the impairment of spatial and contextual fear memories in knock-in mutant mice lacking the T286 autophosphorylation suggest that NMDA receptor-dependent LTP is fundamentally important for hippocampal learning and memory (Giese and Mizuno, 2013). NMDA receptor-dependent LTP results from the depolarisation of synapses and removal of the magnesium block from NMDA receptors, allowing for calcium ions to infiltrate the post-synaptic receptor, resulting in α CaMKII autophosphorylation and neuroplasticity changes (Lisman et al., 2012). Unlike NMDA receptor-dependent LTP, NMDA receptor-independent LTP

occurs more predominantly in aged animals in the hippocampus and involves the activation of L-type voltage-gated calcium channels (VGCC) (Boric et al., 2008), possibly because expression of VGCC increases with age (Oh et al., 2016). Whilst VGCCs can activate α CaMKII autophosphorylation (Pasek et al., 2015), it is unlikely that VGCC-dependent LTP is induced during hippocampus-dependent memory formation in young age (Whitlock et al., 2006; Gruart et al., 2015). However, it is unknown whether in old age VGCC-dependent LTP contributes to memory formation.

Contextual fear conditioning is a hippocampus-dependent memory task that involves the amygdala to form and retain memory, after invoking the mice's fear response in the form of freezing through training with electrical foot shocks (Maren et al., 2013). The tri-synaptic pathway in the hippocampus is critical to memory where input from the entorhinal cortex is processed and filtered to the dentate gyrus, CA3 and then the CA1 brain regions. This predominantly excitatory pathway is modulated by inputs from the amygdala to mediate learning through

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changes in neuronal plasticity (Shechner et al., 2014). The amygdala, which is part of the limbic system plays a role in this emotional learning and is the brain region where connections from sensory cortical inputs are converted into specific autonomic and behavioural responses, such as the fear response demonstrated during contextual fear conditioning (Davis, 1994; Schafe et al., 2005).

Aging impairs contextual fear and spatial memory (Murphy et al., 2004; Peters et al., 2014). A possible explanation for this deterioration may come from age-related difficulties in activating NMDA receptors, raising the threshold for NMDA receptor-dependent LTP induction (Murphy et al., 2004; Bonhaus et al., 1990; Kumar, 2015; Deupree et al., 1993; Moore et al., 1993; Diana et al., 1995; Barnes et al., 1996; Kumar et al., 2007). Furthermore, aging encourages NMDA receptor-independent methods for LTP induction using VGCC channels in the hippocampus (Boric et al., 2008). Moreover, recent studies have shown after complete lesioning of the hippocampus, contextual fear memories can still be formed after multiple training trials (Wiltgen et al., 2006), suggesting that other brain regions such as the amygdala may be involved in learning and memory. Therefore, it is important to study how aging can affect critical molecular mechanisms such as autophosphorylation of α CaMKII after CFC in associated brain regions. We systematically analysed the phosphorylated T286 (active) and total levels of α CaMKII in the hippocampus and amygdala after contextual fear training in young and aged mice.

2. Materials and methods

2.1. Subjects

Experiments were conducted using female C57BL/6J inbred mice (Harlan, NL), aged mice ($n = 10$) were 18 months and young mice ($n = 10$) 3 months old. All work-involving mice were conducted in accordance with the UK Animals Scientific Procedures Act 1986.

2.2. Training

Half of the mice from each age group was randomly allocated to contextual fear conditioning (CFC), creating four unique groups; young trained (YT, $n = 5$), young untrained (YU, $n = 5$), aged trained (AT, $n = 5$) and aged untrained (AU, $n = 5$). Mice dedicated to training were placed one at a time in an enclosed observational chamber (MedAssociates), with their first shock (0.7 mA) lasting 2 s, being administered 148 s after insertion. Subsequent shocks were applied at 90 s intervals and this was repeated four times (five shocks in total), before the mice were returned to their habitat 30 s after their final shock. This conditioning protocol led to similar levels of 24-h contextual fear memory in young and aged mice (Aziz et al., in preparation).

2.3. Tissue preparation

Mice were perfused and brains isolated two hours post-conditioning, brains were post-fixed in 4% paraformaldehyde, 30% sucrose and flash frozen following protocols from previous studies (Radwanska et al., 2011). 40 μ m thick slices were achieved using a cryotome (Leica Biosystems, DE) and slices at -1.50 mm bregma point were isolated and stained following a previous method for phosphorylated α CaMKII

(T286) (Ouyang et al., 1997) and a modified protocol for total α CaMKII dilutions (Cox and Racca, 2013), antibodies shown in Table 1. Antibody specificity was determined by staining of a sample with no primary antibodies. Staining for phosphorylated and total α CaMKII was conducted on adjacent brain samples. Images were acquired for two slices per animal using Axio Imager 2 with Apotome (Zeiss, DE) of the hippocampus (CA3 stratum radiatum), amygdala (basolateral, lateral and central regions) and hippocampus (CA1 stratum oriens) to use as a background.

2.4. Data analysis

Mean α CaMKII density was calculated by taking measurements from three identical rectangular areas on images and taken as a ratio of mean background density using the corresponding mean CA1 stratum oriens α CaMKII value. If mean background density was indeterminate for a particular slice due to damage during preparation, then the background value of the corresponding slice for that animal was used. There were no instances where both background values for the same animal was missing. Ratios after background calculations of the two slices for each brain areas were averaged to give mean density per animal, for both phosphorylated (T286) and total α CaMKII. The proportion of α CaMKII that was phosphorylated compared to the total was calculated as a ratio, by dividing phosphorylated with total values for each animal. An average group value was used to treat missing values and allowed for a ratio for each animal to be obtained. All data was standardised accordingly with YU values by dividing with the corresponding YU data. Graphs were plotted as mean \pm standard error of mean on GraphPad Prism v7.0, US.

2.5. Statistical analysis

Two-way ANOVA was conducted for all animal groups and Tukey post-hoc analysis was undertaken to signify specific differences in total, phosphorylated and ratio of phosphorylated:total α CaMKII, between young and aged mice in different brain regions. For the ratio of phosphorylated:total α CaMKII in CA3 hippocampus, lateral and central amygdala, original data was not normally distributed and instead a two-way ANOVA was conducted further to log transformation. All other outcome variables were found to be normally distributed and of equal variance. Statistical tests were carried out on SigmaPlot v13.0, US.

3. Results

The autophosphorylation (T286) of α CaMKII is essential for NMDA receptor-dependent LTP (Lisman et al., 2012; Giese and Mizuno, 2013), and also occurs after the induction of VGCC-dependent LTP (Pasek et al., 2015), lasting for several hours (Irvine et al., 2006). We tested whether aging and contextual fear conditioning can alter the levels of phosphorylated α CaMKII in hippocampus and amygdala nuclei (Fig. 1), by analysing the T286 phosphorylated (activated) and total levels of α CaMKII. It was previously shown that phosphorylation of α CaMKII occurs two hours after contextual fear conditioning (Atkins et al., 1998). Similarly, in our study we have chosen a two-hour time point to study changes in phosphorylated α CaMKII for varying brain regions.

The total α CaMKII in the CA3 stratum radiatum of the hippocampus

Table 1

Primary and secondary antibodies used for immunohistochemistry staining for phosphorylated and total α CaMKII. Antibody concentrations, name and conditions for staining displayed.

Experiment type	Antibody type	Name	Species	Special Conditions	Dilution
Phosphorylated α CaMKII (T286)	Primary	ABCAM, UK (AB5683)	Rabbit polyclonal	18 h rt	1:300
	Secondary	Life Tech, US (A11034)	Goat anti-Rabbit	2 h rt	1:2000
Total α CaMKII	Primary	Abnova Corp., ROC (MAB8699)	Mouse monoclonal	72 h 4 °C	1:2000
	Secondary	Life Tech, US (A11004)	Goat anti-mouse	2 h rt	1:500

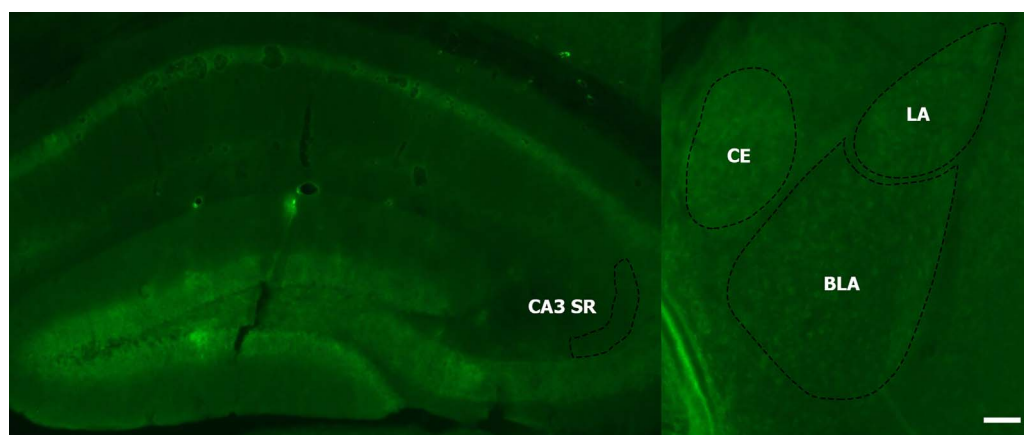


Fig. 1. Low magnification images of the brain regions stained with pCaMKII antibody. Hippocampus (left) showing CA3 stratum radiatum (SR) and amygdala (right) showing lateral (LA), basolateral (BLA) and central (CE) regions as indicated. Images stained using eGFP (green). Scale bar, 40 μ m. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

(Fig. 2A), showed no significant differences by two-way ANOVA for training ($F_{(1,15)} = 0.28$, $p = 0.60$), aging ($F_{(1,15)} = 0.43$, $p = 0.52$) and interaction of training and aging ($F_{(1,15)} = 0.58$, $p = 0.46$) (Fig. 2B). The phosphorylated α CaMKII (Fig. 2A), showed significant differences for training ($F_{(1,11)} = 4.9$, $p = 0.05$) but not for ageing ($F_{(1,11)} = 0.7$, $p = 0.44$) or interaction of training and aging ($F_{(1,11)} = 2.4$, $p = 0.15$) (Fig. 2C). For the ratio of phosphorylated:total α CaMKII, training was significant ($F_{(1,16)} = 5.0$, $p = 0.04$), but not for aging ($F_{(1,16)} = 2.2$, $p = 0.16$) or interaction ($F_{(1,16)} = 3.3$, $p = 0.09$) (Fig. 2D). Tukey's post hoc analysis showed increases to phosphorylated α CaMKII after contextual fear training in young mice ($p = 0.03$) but not aged mice ($p = 0.64$) (Fig. 2C). The ratio of phosphorylated:total α CaMKII after training significantly increased in young mice ($p = 0.01$) but not in aged mice ($p = 0.77$). However, this ratio was significantly increased in aged when compared to young untrained mice ($p = 0.03$) (Fig. 2D) (Table 2).

The total α CaMKII in the lateral amygdala (Fig. 3A), showed significant differences by two-way ANOVA for training ($F_{(1,16)} = 8.3$, $p = 0.01$), but not for aging ($F_{(1,16)} = 2.4$, $p = 0.14$) or interaction of training and aging ($F_{(1,16)} = 3.4$, $p = 0.08$) (Fig. 3B). The phosphorylated α CaMKII (Fig. 3A), showed no significant differences for training ($F_{(1,13)} = 0.2$, $p = 0.64$), ageing ($F_{(1,13)} = 0.3$, $p = 0.60$), and interaction of training and aging ($F_{(1,13)} = 0.3$, $p = 0.59$) (Fig. 3C). For the ratio of phosphorylated:total α CaMKII, there was also no significant effect of training ($F_{(1,16)} = 3.9$, $p = 0.07$), ageing ($F_{(1,16)} = 1.1$, $p = 0.30$) and interaction ($F_{(1,16)} = 1.2$, $p = 0.28$) (Fig. 3D) (Table 2). The non-significant trend for an up-regulation of phosphorylated:total α CaMKII after conditioning in young mice is consistent with the detection of an up-regulation, using immuno-electronmicroscopy (Rodrigues et al., 2004).

The total α CaMKII in the central amygdala (Fig. 4A), showed no significant differences by two-way ANOVA for training ($F_{(1,16)} = 3.5$,

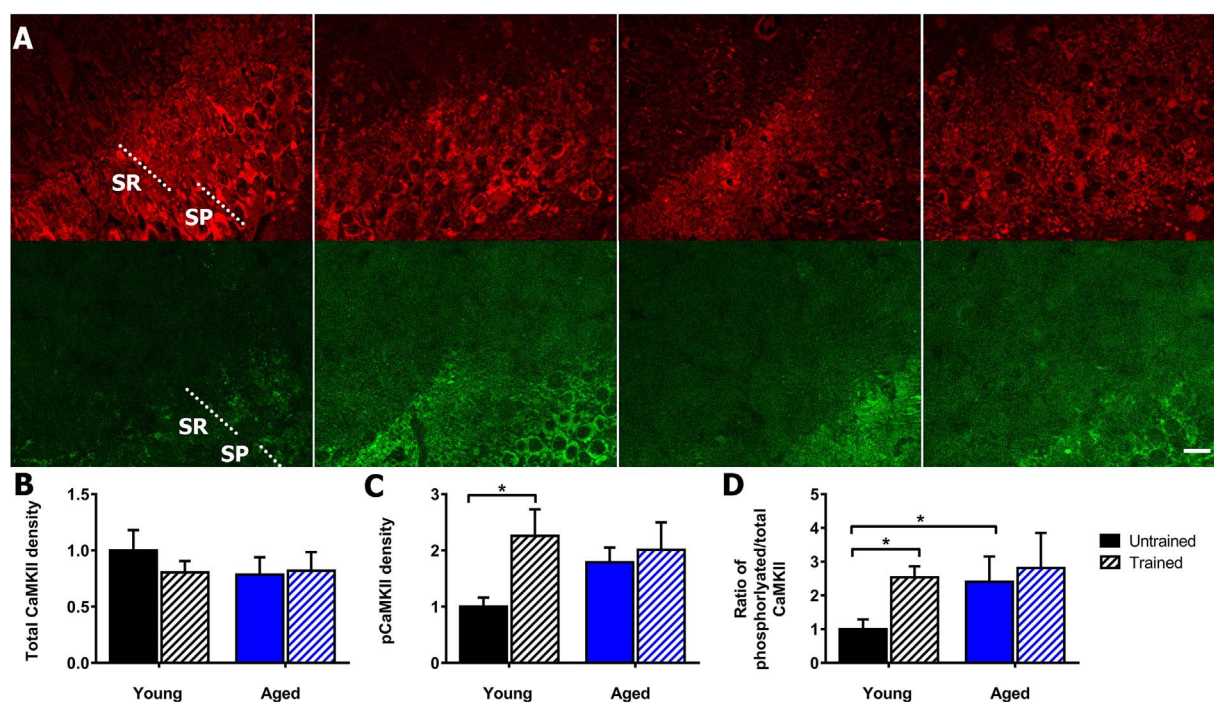


Fig. 2. Age-dependent impairment of α CaMKII activation in stratum radiatum of hippocampal area CA3 after contextual fear conditioning. A) Representative images of phosphorylated and total α CaMKII in the stratum radiatum (SR) and stratum pyramidal (SP) of the CA3 hippocampus. Total α CaMKII stained using dsRed (red), phosphorylated α CaMKII stained using eGFP (green). Young untrained (far left), young trained (centre left), aged untrained (centre right), aged trained (far right), scale 20 μ m. B) Contextual fear conditioning does not alter total α CaMKII levels in the stratum radiatum of the CA3 hippocampus in young and aged mice. C) Contextual fear conditioning induced an increase in autophosphorylation of α CaMKII in young mice ($p = 0.025$), but there was no significant change in aged mice. D) Contextual fear conditioning induced an increase in the ratio of autophosphorylated to total α CaMKII in young ($p = 0.012$), but not aged mice. Ageing induced an increase in ratio of phosphorylated:total α CaMKII in untrained mice ($p = 0.033$). Mean \pm standard error of mean, * ($p < 0.05$). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

Table 2

Mean and S.E.M. of total, phosphorylated and phosphorylated:total α CaMKII values for young untrained (YU), young trained (YT), aged untrained (AU) and aged trained (AT) mice groups in the CA3 hippocampus, lateral (LA), central (CE) and basolateral (BLA) amygdala.

	Mean Total (S.E.M.)				Mean Phosphorylated (S.E.M.)				Mean Phosphorylated:Total (S.E.M.)			
	YU	YT	AU	AT	YU	YT	AU	AT	YU	YT	AU	AT
CA3	1.0 (0.2)	0.8 (0.1)	0.8 (0.2)	0.8 (0.2)	1.0 (0.2)	2.2 (0.5)	1.8 (0.3)	2.0 (0.5)	1.0 (0.3)	2.5 (0.3)	2.4 (0.7)	2.8 (1.0)
LA	1.0 (0.1)	0.5 (0.0)	0.7 (0.1)	0.5 (0.1)	1.0 (0.2)	1.3 (0.2)	1.4 (0.6)	1.3 (0.3)	1.0 (0.3)	2.3 (0.3)	2.4 (1.1)	2.5 (0.8)
CE	1.0 (0.2)	0.7 (0.1)	0.8 (0.1)	0.7 (0.1)	1.0 (0.2)	1.1 (0.5)	1.5 (0.5)	1.1 (0.1)	1.0 (0.3)	1.6 (0.5)	1.6 (0.5)	1.3 (0.2)
BLA	1.0 (0.1)	0.8 (0.1)	0.9 (0.1)	0.7 (0.0)	1.0 (0.2)	1.0 (0.1)	1.0 (0.1)	1.1 (0.1)	1.0 (0.3)	1.1 (0.2)	1.1 (0.2)	1.3 (0.1)

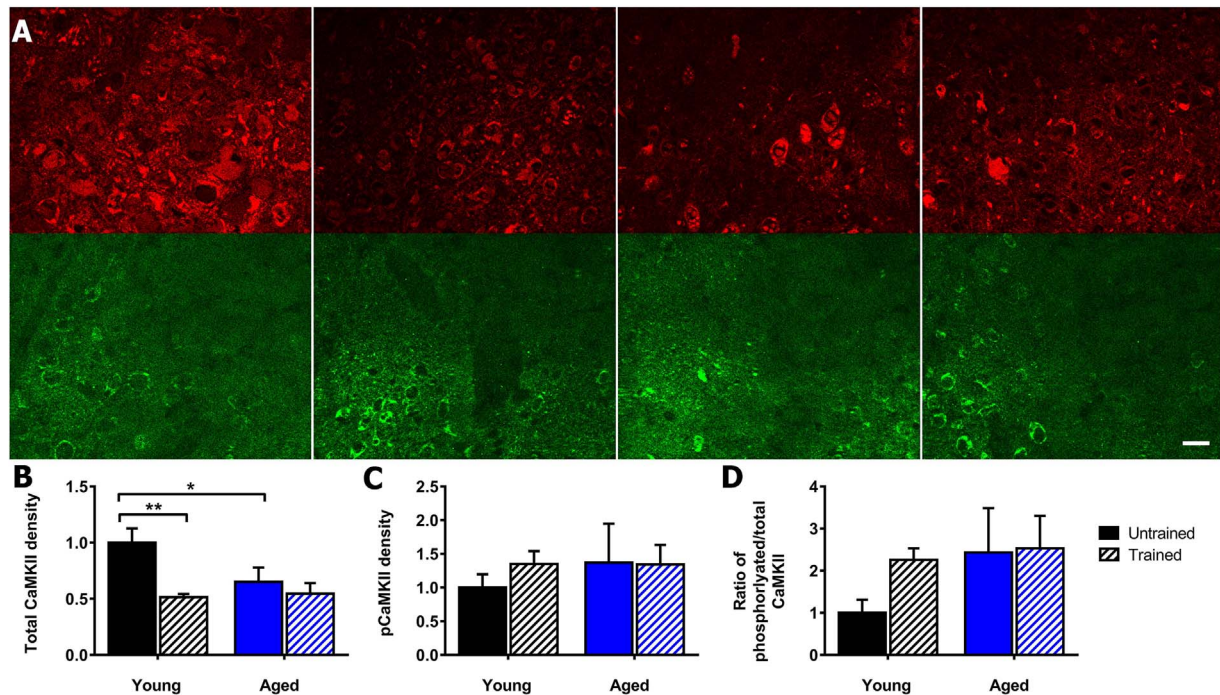


Fig. 3. α CaMKII autophosphorylation in lateral amygdala in young and aged mice after contextual fear conditioning. A) Representative images of phosphorylated and total α CaMKII in the lateral amygdala. Total α CaMKII stained using dsRed (red), phosphorylated α CaMKII stained using eGFP (green). Young untrained (far left), young trained (centre left), aged untrained (centre right), aged trained (far right), scale 20 μ m. B) Contextual fear conditioning significantly decreased total α CaMKII levels in the lateral amygdala in young ($p = 0.004$) but not aged mice. Ageing reduced the levels of total α CaMKII in untrained mice ($p = 0.029$). C) Contextual fear conditioning does not alter α CaMKII autophosphorylation in young and aged mice. D) Contextual fear conditioning does not alter the ratio of phosphorylated:total α CaMKII in the lateral amygdala in young and aged mice, although there was a non-significant trend of a conditioning-induced up-regulation in young age. Mean \pm standard error of mean. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

$p = 0.08$), aging ($F_{(1,16)} = 0.2$, $p = 0.65$) and interaction of training and aging ($F_{(1,16)} = 0.7$, $p = 0.42$) (Fig. 4B). The phosphorylated α CaMKII (Fig. 4A), showed no significant differences for training ($F_{(1,11)} = 0.1$, $p = 0.72$), aging ($F_{(1,11)} = 0.4$, $p = 0.54$) and interaction of training and aging ($F_{(1,11)} = 0.6$, $p = 0.44$) (Fig. 4C) (Table 2). For the ratio of phosphorylated:total α CaMKII, there was also no significant effect of training ($F_{(1,16)} = 0.6$, $p = 0.44$), ageing ($F_{(1,16)} = 0.7$, $p = 0.41$) and interaction ($F_{(1,16)} = 1.2$, $p = 0.29$) (Fig. 4D) (Table 2).

The total α CaMKII in the basolateral amygdala (Fig. 5A), showed no significant differences by two-way ANOVA for training ($F_{(1,16)} = 2.4$, $p = 0.14$), aging ($F_{(1,16)} = 0.9$, $p = 0.35$) and interaction of training and aging ($F_{(1,16)} < 0.1$, $p = 0.97$) (Fig. 5B). The phosphorylated α CaMKII (Fig. 5A), showed no significant differences for training ($F_{(1,13)} = 0.02$, $p = 0.90$), ageing ($F_{(1,13)} = 0.09$, $p = 0.76$) and interaction of training and aging ($F_{(1,13)} < 0.001$, $p = 0.99$) (Fig. 5C). For the ratio of phosphorylated:total α CaMKII, there was also no significant effect of training ($F_{(1,13)} = 0.39$, $p = 0.54$), ageing ($F_{(1,16)} = 0.4$, $p = 0.55$) and interaction ($F_{(1,13)} = 0.01$, $p = 0.99$) (Fig. 5D) (Table 2).

4. Discussion

Experience-dependent molecular changes underlying synaptic plasticity in learning and memory can be modified by aging. An accurate and detailed understanding of how these molecules play a crucial role in hippocampus-dependent memory tasks and how they are affected by aging are poorly understood. We investigated this issue by looking at levels of a critical molecular step in induction of LTP, α CaMKII autophosphorylation at T286, which persists for some time after LTP induction (Irvine et al., 2006), in the hippocampus and amygdala in young and aged mice after contextual fear conditioning. For these experiments we compared the levels of autophosphorylation of α CaMKII and the ratio of phosphorylated:total α CaMKII. We used the ratio of phosphorylated:total α CaMKII to demonstrate the proportion of available α CaMKII that was autophosphorylated. As this would serve as a good marker in determining any increases in protein activity at any one point in time, considering any changes to the level of total α CaMKII after training or aging. It might also be a proxy for calcium signalling (Pasek et al., 2015).

Our experiments showed that contextual fear conditioning increases the levels of both phosphorylated and the ratio of phosphorylated:total

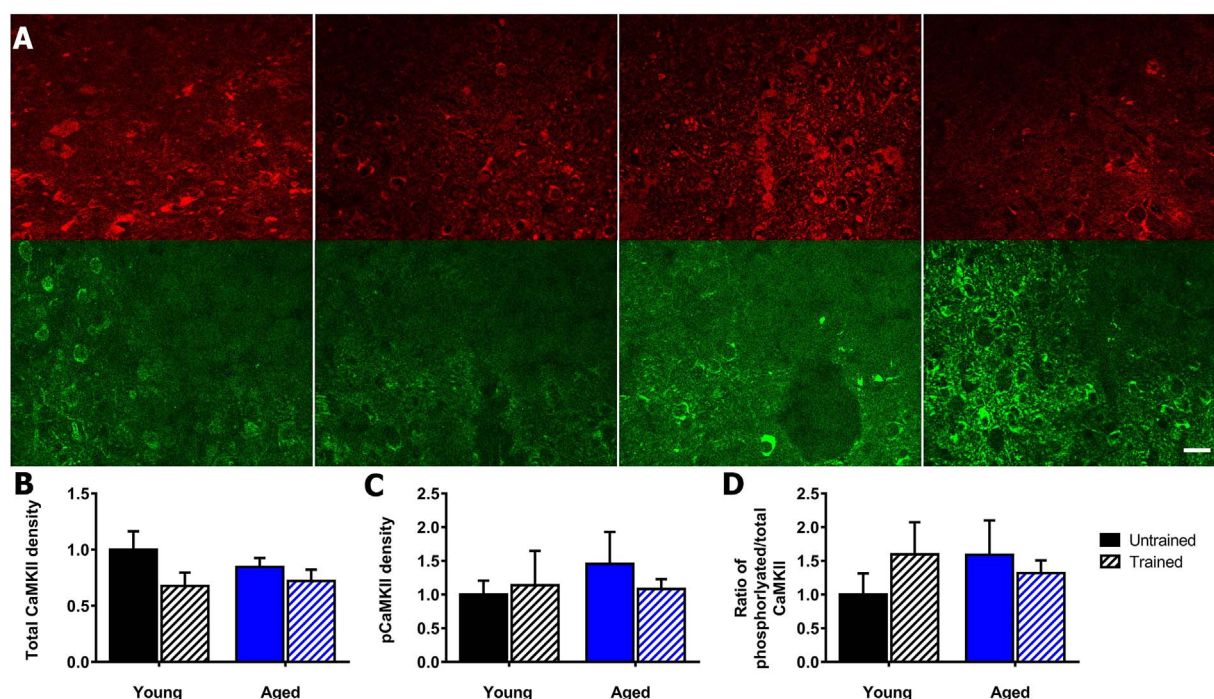


Fig. 4. α CaMKII autophosphorylation in central amygdala in young and aged mice after contextual fear conditioning. A) Representative images of phosphorylated and total α CaMKII in the central amygdala. Total α CaMKII stained using dsRed (red), phosphorylated α CaMKII stained using eGFP (green). Young untrained (far left), young trained (centre left), aged untrained (centre right), aged trained (far right), scale 20 μ m. B) Contextual fear conditioning does not alter total α CaMKII levels in the central amygdala in young and aged mice. C) Contextual fear conditioning does not alter α CaMKII autophosphorylation in young and aged mice. D) Contextual fear conditioning does not alter the ratio of phosphorylated:total α CaMKII in the central amygdala in young and aged mice. Mean \pm standard error of mean. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

α CaMKII in the hippocampal CA3 stratum radiatum in young, but not aged mice. To the best of our knowledge, this finding represents the first experiment conducted to determine the changes to α CaMKII

autophosphorylation in the CA3 region after CFC. Previous studies have shown by blocking autophosphorylation of α CaMKII, contextual fear and spatial memories are impaired in mice (Irvine et al., 2005; Need

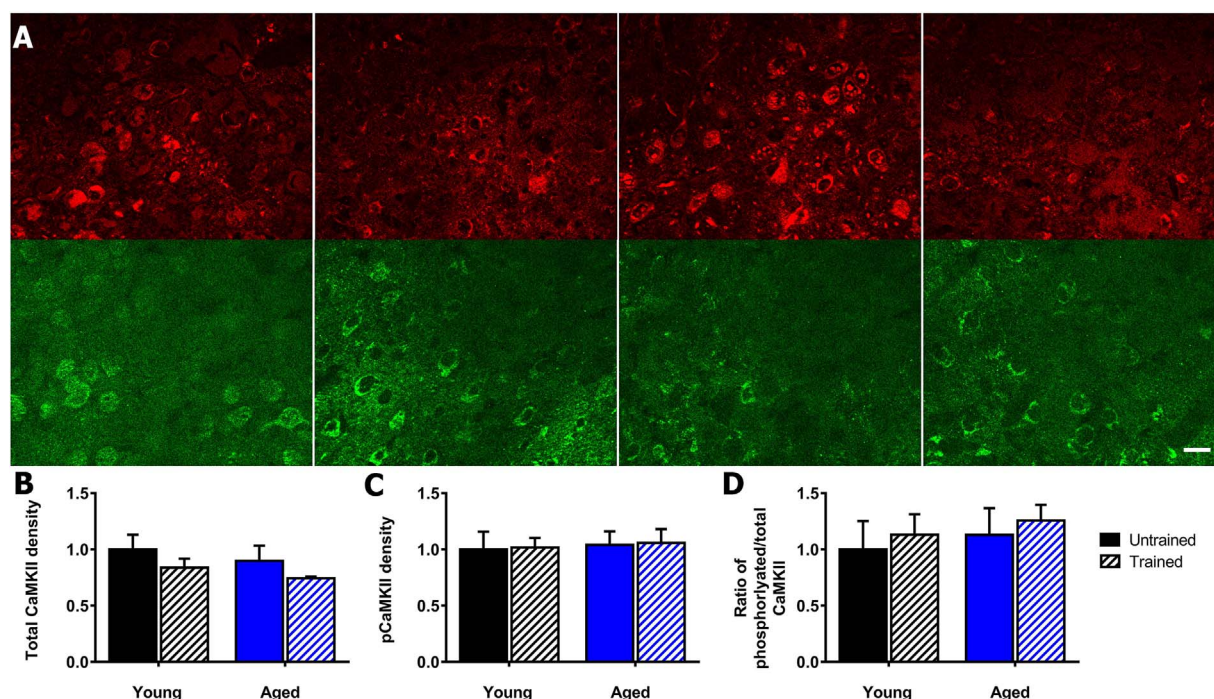


Fig. 5. α CaMKII autophosphorylation in basolateral amygdala in young and aged mice after contextual fear conditioning. A) Representative images of phosphorylated and total α CaMKII in the basolateral amygdala. Total α CaMKII stained using dsRed (red), phosphorylated α CaMKII stained using eGFP (green). Young untrained (far left), young trained (centre left), aged untrained (centre right), aged trained (far right), scale 20 μ m. B) Contextual fear conditioning does not alter total α CaMKII levels in the basolateral amygdala in young and aged mice. C) Contextual fear conditioning does not alter α CaMKII autophosphorylation in the basolateral amygdala in young and aged mice. D) Contextual fear conditioning does not alter the ratio of phosphorylated:total α CaMKII in the basolateral amygdala in young and aged mice. Mean \pm standard error of mean. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

and Giese, 2003). Our study suggests that an increase in phosphorylated α CaMKII after training in the CA3 region of the hippocampus may contribute to contextual fear memory formation in young age. Thus, in young mice the memory mechanism in the CA3 region after training might be associated with NMDA receptor-dependent LTP as suggested in a previous trace eyeblink conditioning study (Gruart et al., 2015).

Surprisingly, we found that in young, but not aged mice, total α CaMKII levels decreased in the lateral amygdala after CFC. The mechanism underlying this decreased expression is unclear. Interestingly, the decrease in α CaMKII levels is associated with a trend of an increase in phosphorylated:total α CaMKII in young mice, which is consistent with the detection of an up-regulation, using immuno-electronmicroscopy (Rodrigues et al., 2004). Our results indicate that this up-regulation does not occur in old age.

Aged mice which underwent CFC training, showed no differences for both levels of phosphorylated and total α CaMKII in the hippocampus and amygdala. This lack of change could be attributed to age-related difficulties in the induction and maintenance of LTP through an increased threshold for induction and difficulties in maintaining synaptic plasticity (Murphy et al., 2004; Bonhaus et al., 1990; Kumar, 2015; Deupree et al., 1993; Moore et al., 1993; Diana et al., 1995; Barnes et al., 1996; Kumar et al., 2007). Therefore, in line with our expectations there were no difference in phosphorylated α CaMKII and the ratio of autophosphorylation to total levels of α CaMKII after training in aged mice for all brain regions examined. This finding also suggests that it would be unlikely for VGCC-dependent LTP to occur after contextual fear conditioning in aged mice, as this would otherwise lead to an increase in the level of phosphorylated α CaMKII.

Interestingly, we found that for untrained mice, aging increased the ratio of phosphorylated:total α CaMKII in the CA3 stratum radiatum. This increase in baseline autophosphorylation may be due to the well-known increase in calcium entry through VGCC (Oh et al., 2016). The effect of such raised calcium levels in aged mice could be even further amplified by the depolarisation of post-synaptic spines which would lead to NMDA receptor activation and increased calcium influx, causing more autophosphorylation of α CaMKII. This persistent calcium elevation, could be a reason as to why training does not increase phosphorylated α CaMKII any further in aged mice, where a ceiling effect may have been reached. This is when the maximal level of phosphorylation at any point in time is reached and therefore any extra training-induced stimuli will not see any further increase to autophosphorylation.

In summary, our findings suggest that the hippocampus makes a unique contribution to contextual fear memory formation. Our results lead us to suggest that in young, but not old age NMDA receptor-dependent LTP in hippocampal area CA3 may contribute to contextual fear memory formation.

Declaration of interest

Authors declare no conflict of interest.

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References

Atkins, C.M., Selcher, J.C., Petraitis, J.J., Trzaskos, J.M., Sweatt, D., 1998. The MAPK

- cascade is required for mammalian associative learning. *Nat. Neurosci.* 1, 602–609.
- Barnes, C.A., Rao, G., McNaughton, B.L., 1996. Functional integrity of NMDA-dependent LTP induction mechanisms across the lifespan of F-344 rats. *Learn. Mem.* 3, 124–137.
- Bonhaus, D.W., Perry, W.B., McNamara, J.O., 1990. Decreased density but not number, of N-methyl-D-aspartate, glycine and phencyclidine binding sites in hippocampus of senescent rats. *Brain Res.* 532, 82–86.
- Boric, K., Munoz, P., Gallagher, M., Kirkwood, A., 2008. Potential adaptive function for altered long-term potentiation mechanisms in aging hippocampus. *J. Neurosci.* 28, 8034–8039.
- Cox, D.J., Racca, C., 2013. Differential dendritic targeting of AMPA receptor subunit mRNAs in adult rat hippocampal principal neurons and interneurons. *J. Comp. Neurol.* 521, 1954–2007.
- Davis, M., 1994. The role of the amygdala in emotional learning. *Int. Rev. Neurobiol.* 36, 225–266.
- Deupree, D.L., Bradley, J., Turner, D.A., 1993. Age-related alterations in potentiation in the CA1 region in F344 rats. *Neurobiol. Aging* 14, 249–258.
- Diana, G., Domenici, M.R., Scotti de Carolis, A., Loizzo, A., Sagratella, S., 1995. Reduced hippocampal CA1 Ca^{2+} -induced long-term potentiation is associated with age-dependent impairment of spatial learning. *Brain Res.* 686, 107–110.
- Giese, K.P., Mizuno, K., 2013. The roles of protein kinases in learning and memory. *Learn. Mem.* 16, 540–552.
- Giese, K.P., 2012. Long-term potentiation and memory. In: Giese, K.P. (Ed.), *Memory Mechanisms in Health and Disease*. World Scientific (Singapore), pp. 1–17.
- Gruart, A., Leal-Campanario, R., Lopez-Ramos, J.C., Delgado-García, J.M., 2015. Functional basis of associative learning and its relationships with long-term potentiation evoked in the involved neural circuits: lessons from studies in behaving mammals. *Neurobiol. Learn. Mem.* 124, 3–18.
- Irvine, E.E., Vernon, J., Giese, K.P., 2005. AlphaCaMKII autophosphorylation contributes to rapid learning but is not necessary for memory. *Nat. Neurosci.* 8, 411–412.
- Irvine, E.E., von Hertzen, L.S., Plattner, F., Giese, K.P., 2006. alphaCaMKII autophosphorylation: a fast track to memory. *Trends Neurosci.* 29, 459–465.
- Kumar, A., Thinschmidt, J.S., Foster, T.C., King, M.A., 2007. Aging effects on the limits and stability of long-term synaptic potentiation and depression in rat hippocampal area CA1. *J. Neurophysiol.* 98, 594–601.
- Kumar, A., 2015. NMDA receptor function during senescence: implication on cognitive performance. *Front. Neurosci.* 9, 473.
- Lisman, J., Yasuda, R., Raghavachari, S., 2012. Mechanisms of CaMKII action in long-term potentiation. *Nat. Rev. Neurosci.* 13, 175–190.
- Maren, S., Phan, K.L., Liberzon, I., 2013. The contextual brain: implications for fear conditioning: extinction and psychopathology. *Nat. Rev. Neurosci.* 14, 417–428.
- Moore, C.I., Browning, M.D., Rose, G.M., 1993. Hippocampal plasticity induced by primed burst, but not long-term potentiation, stimulation is impaired in area CA1 of aged Fischer 344 rats. *Hippocampus* 3, 57–66.
- Murphy, G.G., Fedorov, N.B., Giese, K.P., Ohno, M., Friedman, E., Chen, R., Silva, A.J., 2004. Increased neuronal excitability synaptic plasticity, and learning in aged Kvbeta1.1 knockout mice. *Curr. Biol.* 14, 1907–1915.
- Need, A.C., Giese, K.P., 2003. Handling and environmental enrichment do not rescue learning and memory impairments in alphaCaMKII(T286A) mutant mice. *Genes Brain Behav.* 2, 132–139.
- Oh, M.M., Simkin, D., Disterhoft, J.F., 2016. Intrinsic hippocampal excitability changes of opposite signs and different origins in CA1 and CA3 pyramidal neurons underlie aging-related cognitive deficits. *Front. Syst. Neurosci.* 10, 52.
- Ouyang, Y., Kantor, D., Harris, K.M., Schuman, E.M., Kennedy, M.B., 1997. Visualization of the distribution of autophosphorylated calcium/calmodulin-dependent protein kinase II after tetanic stimulation in the CA1 area of the hippocampus. *J. Neurosci.* 17, 5416–5427.
- Pasek, J.G., Wang, X., Colbran, R.J., 2015. Differential CaMKII regulation by voltage-gated calcium channels in the striatum. *Mol. Cell. Neurosci.* 68, 234–243.
- Peters, M., Bletsch, M., Stanley, J., Wheeler, D., Scott, R., Tully, T., 2014. The PDE4 inhibitor HT-0712 improves hippocampus-dependent memory in aged mice. *Neuropsychopharmacology* 39, 2938–2948.
- Radwanska, K., Medvedev, N.I., Pereira, G.S., Engmann, O., Thiede, N., Moraes, M.F., et al., 2011. Mechanism for long-term memory formation when synaptic strengthening is impaired. *Proc. Natl. Acad. Sci. U. S. A.* 108, 18471–18475.
- Rodrigues, S.M., Farb, C.R., Bauer, E.P., LeDoux, J.E., Schafe, G.E., 2004. Pavlovian fear conditioning regulates Thr286 autophosphorylation of calcium/calmodulin-dependent protein kinase II at lateral amygdala synapses. *J. Neurosci.* 24, 3281–3288.
- Schafe, G.E., Doyere, V., LeDoux, J.E., 2005. Tracking the fear engram: the lateral amygdala is an essential locus of fear memory storage. *J. Neurosci.* 25, 10010–10014.
- Shechner, T., Hong, M., Britton, J.C., Pine, D.S., Fox, N.A., 2014. Fear conditioning and extinction across development: evidence from human studies and animal models. *Biol. Psychol.* 100, 1–12.
- Whitlock, J.R., Heynen, A.J., Epstein, M.H., Carpenter, R.L., Bear, M.F., 2006. Learning induces long-term potentiation in the hippocampus. *Science* 313, 1093–1097.
- Wiltgen, B.J., Sanders, M.J., Anagnostaras, S.G., Sage, J.R., 2006. Fanselow MS. Context fear learning in the absence of the hippocampus. *J. Neurosci.* 26, 5484–5491.